

END-CAPPED QUINOXALINE-CONTAINING HYPERBRANCHED
AROMATIC POLY(ETHER-KETONES)

CROSS REFERENCE TO RELATED APPLICATION

5 This application is a continuation-in-part of application serial number 10/695,735, filed 10/23/2003, which in turn claims benefit of priority of the filing date of Provisional Application serial number 60/453,334, filed 02-28-2003.

RIGHTS OF THE GOVERNMENT

The invention described herein may be manufactured and used by or for the Government of the United States for all governmental purposes without the payment of any royalty.

BACKGROUND OF THE INVENTION

The present invention relates to a new quinoxaline-containing hyperbranched ether-ketone polymers.

Dendritic macromolecules such as dendrimers and hyperbranched polymers are

5 a new class of highly branched polymers that have distinctly different properties from their linear analogs. Both dendrimers and hyperbranched polymers have much lower solution and melt viscosities than their linear analogs of similar molecular weights. They also have a large number of chain-ends whose collective influence dictates their overall physical and/or chemical behaviors. These features are attractive in terms of

10 processability and offering flexibility in engineering required properties for specific applications. However, there is a practical advantage that hyperbranched polymers have over dendrimers at “raw material” level. Although dendrimers have precisely controlled structures (designated as generations), their preparations generally involve tedious, multi-step sequences that are impractical and costly in scale-up production.

15 Synthesis of a hyperbranched polymer, on the other hand, is a one-pot process. Large quantities of hyperbranched polymers can be easily produced from AB_x ($x \geq 2$) monomers.

Because of their excellent thermal and mechanical properties, as well as their optical and electronic characteristics, aromatic, fused heterocyclic polymers such as

20 polyquinoxalines and polybenzoxazoles continue to attract considerable attention. However, they have limited processability due to the nature of fused ring systems. Their insolubility and their softening temperatures are generally above their degradation temperatures. Chemical modification on the these materials, for example, by the use of

solubilizing pendants or flexible units in the main chain, has been successful to improve their processability, allowing the optimization of their properties as a function of processability. Another viable approach to achieving this objective is to incorporate the elements of local rigidity and global randomness into the macromolecular architecture.

- 5 Local rigidity provides the thermal, electronic and optical characteristics of the aromatic fused systems while global randomness frustrates entanglement of the polymer chains, leading to greater solubility. Dendritic structures clearly embody these qualities. However, as noted previously, hyperbranched structures have greater synthetic practicality.

10 Accordingly, it is an object of the present invention to provide novel quinoxaline-containing hyperbranched poly(ether-ketones).

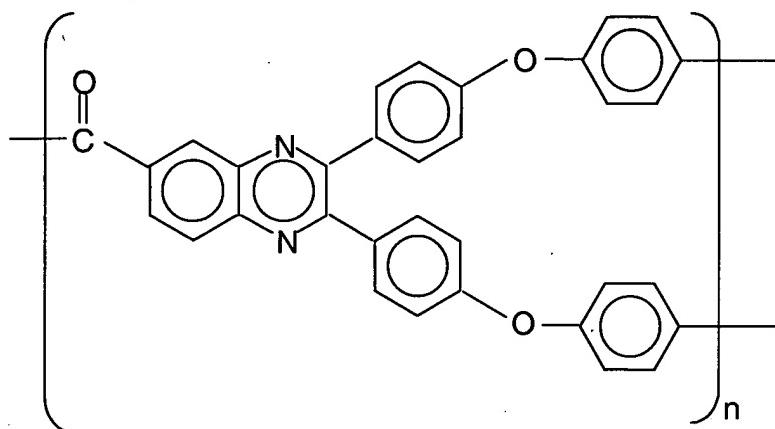
It is another object of the present invention to provide novel end-capped quinoxaline-containing hyperbranched poly(ether-ketones).

15 It is another object of the present invention to provide a novel method for preparing end-capped quinoxaline-containing hyperbranched poly(ether-ketones).

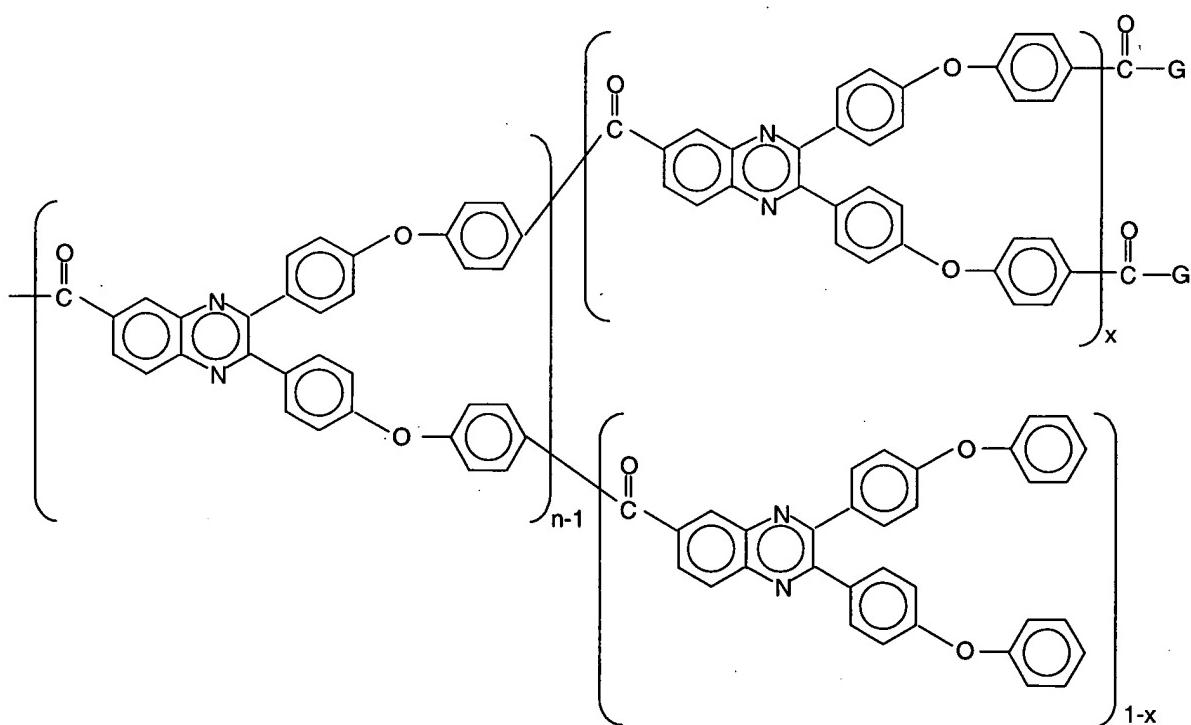
Other objects and advantages of the invention will be set forth in part in the description which follows, and in part will be obvious from the description, or may be learned by practice of the invention. The objects and advantages of the invention may be realized and attained by means of the instrumentalities and combinations particularly 20 pointed out in the appended claims.

SUMMARY OF THE INVENTION

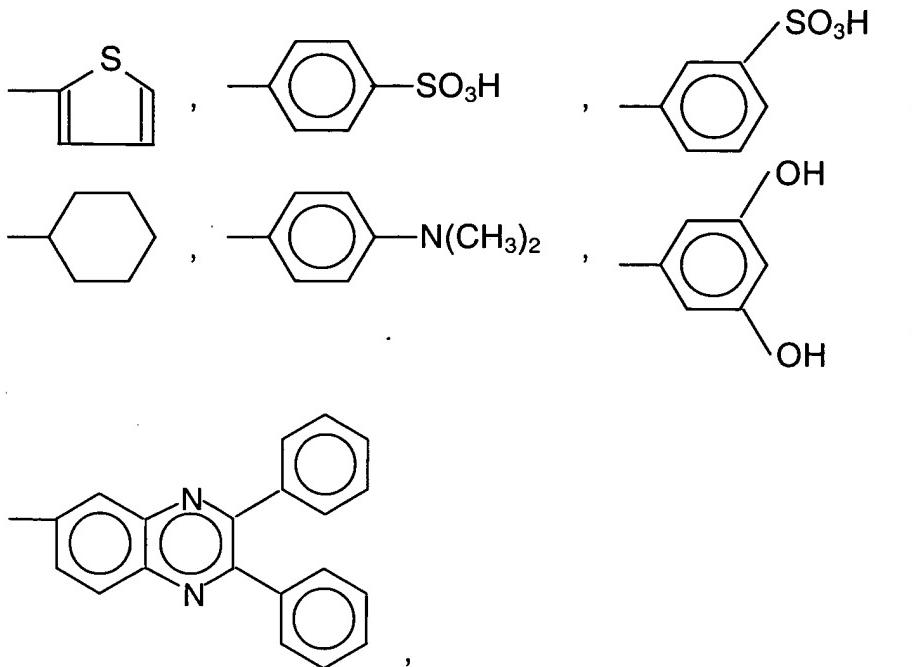
In accordance with the present invention there is provided a quinoxaline-containing hyperbranched ether-ketone polymer having repeating units of the formula:



5 Also provided is an end-capped quinoxaline-containing hyperbranched ether-ketone polymer having repeating units of the formula:



wherein n represents the degree of polymerization for the parent hyperbranched polymer, x represents the degree of endgroup functionalization, x has a value of 0.05 to 1.0, and G is selected from the group consisting of

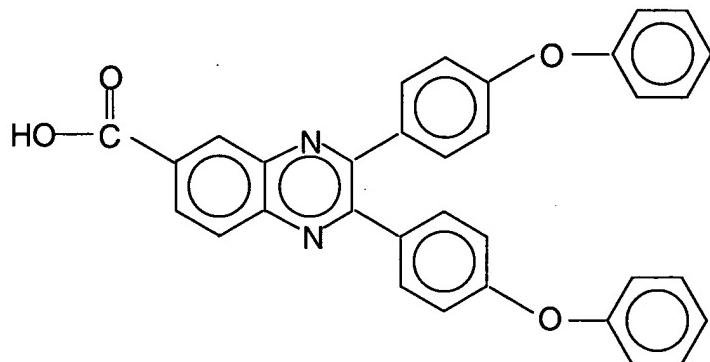


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and aliphatic carboxylic acids having 2 to 16 carbon atoms, i.e., $\text{C}_q\text{H}_{2q+1}\text{COOH}$, wherein q has a value of 1 to 15.

DETAILED DESCRIPTION OF THE INVENTION

The quinoxaline-containing hyperbranched ether-ketone polymer of this invention is prepared by polymerization of the corresponding AB₂ monomer



5 Preparation of the AB₂ monomer, 2,3-bis(4-phenyloxyphenyl)-6-quinoxaline carboxylic acid, is described in co-pending application serial number _____, filed of even date herewith.

Polymerization of the AB₂ monomer can be conducted in polyphosphoric acid (PPA) at a polymer concentration of about 5 weight percent at a temperature of about 10 130°C. Preliminarily it is helpful to describe the chemistry of phosphoric acids and strong phosphoric acids or polyphosphoric acids as follows: As used herein the term "phosphoric acid(s)" means commercial phosphoric acid(s) containing 85-86% H₃PO₄. The strong phosphoric acids, or polyphosphoric acids referred to as PPA (polyphosphoric acid) are members of a continuous series of amorphous condensed 15 phosphoric acid mixtures given by the formula



or



where the value of n depends on the molar ratio of water to phosphorus pentoxide present.

- In its most general definition, polyphosphoric acid composition can range from distributions where the average value of n is less than unity, giving rise to a mobile liquid, to high values of n, where the polyphosphoric acid is a glass at normal temperatures. Because the species of polyphosphoric acid are in a mobile equilibrium, a given equilibrium composition can be prepared in many ways. For instance, the same distribution or polyphosphoric acid composition could be prepared by either starting with concentrated orthophosphoric acid (H_3PO_4 , n=1) and driving off water or by starting with phosphorus pentoxide (P_2O_5) and adding an appropriate amount of water.

All polyphosphoric acid compositions can be described as a ratio of P_2O_5 and water by reducing the various species present (on paper) to P_2O_5 and water. We will then use the convention that polyphosphoric acid composition will be expressed in terms of a P_2O_5 content (as a percentage) defined as P_2O_5 content

$$15 \quad = (\text{weight of } P_2O_5) / (\text{weight of } P_2O_5 + \text{weight of water}) \times 100.$$

Thus, the P_2O_5 content of pure orthophosphoric acid could be derived by reducing one mole of H_3PO_4 to 0.5 moles P_2O_5 + 1.5 moles H_2O . Converting to weights gives the P_2O_5 content as

$$(0.5 * 142) / ((0.5 * 142) + (1.5 * 18.01)) = 72.4\%$$

20 Similarly, the P_2O_5 content of commercial polyphosphoric acid can be derived in the following way. Polyphosphoric acid is available commercially in two grades, 105% and 115%. These percentages refer to H_3PO_4 content, which means that 100 g of the

two grades contain 105 and 115 grams of H₃PO₄. The P₂O₅ content of 115% polyphosphoric acid can then be calculated knowing the P₂O₅ content of 100% H₃PO₄.

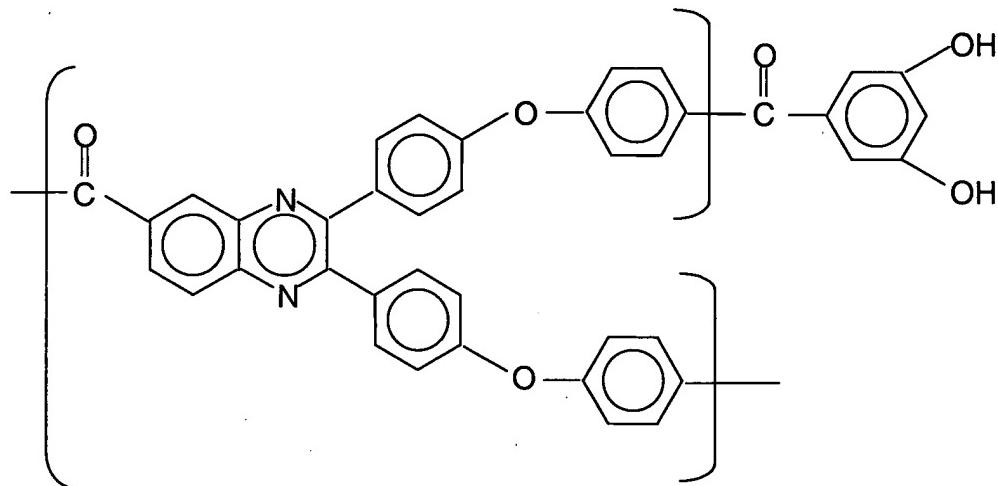
$$(115 * 0.724) / 100 = 83.3\%$$

- We have found that the rate of polymerization can be accelerated by adding
- 5 about 25% additional phosphorus pentoxide (relative to the weight of PPA) to the polymerization mixture, as shown in the examples which follow.

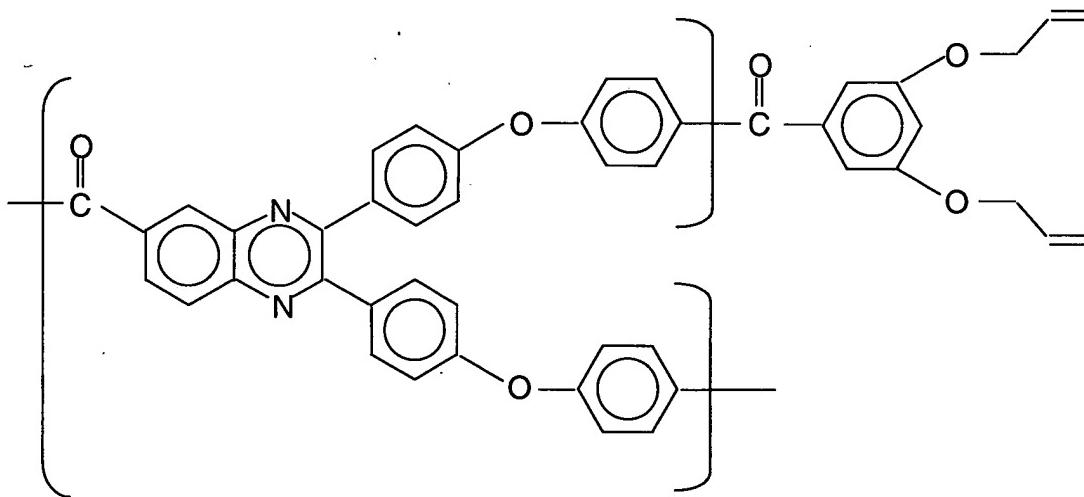
Due to the availability of large number of end-groups, the end-functionalization of hyperbranched polymers can be utilized to tailor their physical properties for various applications. The number of reactive end-groups is equal to the degree of

10 polymerization plus one (DP+1). A sufficient quantity of the selected endcapping compound is added to the polymerization mixture to provide about 5 to 99 mol% endcapping. The hyperbranched polymers can be endcapped using a variety of endcapping compounds, for example, but not limited to, 3,5-dihydroxybenzoic acid, 3-sulfobenzoic acid, 4-sulfobenzoic acid and 2,3-diphenylquinoxaline-6-carboxylic acid. 2-

15 thiophenecarboxylic acid, 4-dimethylaminobenzoic acid, cyclohexanecarboxylic acid, acetic acid through and including palmitic acid, and the like. For example, endcapping the hyperbranched polymer with 3,5-dihydroxybenzoic acid provides 3,5-dihydroxyphenylcarbonyl moieties as chain-end units



which, in turn, can be modified with allylbromide to provide the diallyl-terminated hyperbranched polymer having 3,5-di-allyloxyphenylcarbonyl moieties as chain-end units:



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Other reactive endgroups, such as propargyl and glycidyl functions, can be prepared from the 1,3-dihydroxyphenyl-terminated derivative using similar reactions.

The hyperbranched polymers of this invention are suitable for use in applications where the material will be subject to high service temperatures.

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The following examples illustrate the invention:

Example 1

Polymerization of 2,3-bis(4-phenyloxyphenyl)-6-quinoxaline-carboxylic acid (AB₂ monomer) in PPA at 130°C

- 5 Into 250 mL resin flask equipped with a high-torque mechanical stirrer and nitrogen inlet and outlet, a pressure regulator, and an addition port, polyphosphoric acid (PPA, 60g) was charged. The PPA was degassed under reduced pressure by freezing in liquid nitrogen and melting in warm water several times. Then the monomer, 2,3-bis(4-phenyloxyphenyl)-6-quinoxaline-carboxylic acid (3.0g, 5.9 mmol), was introduced.
- 10 As soon as the monomer was added and the stirring started, the mixture became deep blue-purple. The mixture was heated to 130°C and kept at this temperature for 48h. Although the mixture had become deep red, its viscosity was not significantly increased. The reaction mixture was allowed to cool to 60-70°C and water was added to precipitate the polymer. The resulting mixture was warm at 60-70°C overnight under the nitrogen.
- 15 The resulting bright yellow solids were collected by suction filtration, washed with diluted ammonium hydroxide, and large amount of water. The polymer was finally dried under reduced pressure (0.05 mmHg) at 200°C for 150h to give essentially quantitative yield:
[η]=0.07 dL/g (0.5% solution in MSA at 30.0±0.1°C). Anal. Calcd. for C₃₃H₂₀N₂O₃ C, 80.47%; H, 4.09%; N, 5.69%; O, 9.75%. Found: C, 78.69%; H, 4.34%; N, 5.27%; O, 20 10.81%.

Example 2

Polymerization of AB₂ monomer in PPA/P₂O₅ at 130°C

- Into a 250 mL resin flask equipped with a high-torque mechanical stirrer, nitrogen inlet and outlet, a pressure regulator and an addition port, PPA (83% assay, 80g) was placed and stirred with dried nitrogen purging at 100°C for 10h. The monomer 2,3-

bis(4-phenoxyphenyl)-6-quinoxaline-carboxylic acid (4.0g) was added and heated to 130°C until it became a homogeneous mixture. It usually took about 1 h. The color of mixture became dark brown. P₂O₅ (20.0g ; 25 wt% relative to PPA used) was then added in one portion and the temperature was maintained at 130°C for 24h. The 5 mixture became very viscous after 2h at 130°C and started to stick to the stirring rod. At the end of the reaction, water was added into the flask. The resulting precipitates were collected by suction filtration, washed with diluted ammonium hydroxide followed by a large amounts of water, stirred in boiling water for 100h, and finally dried in the presence of phosphorous pentoxide under reduced pressure (1 mmHg) at 200°C for 10 48h. The yield was essentially quantitative (>99% yield). Its intrinsic viscosity is 0.56 dL/g (MSA, 30 ± 0.1°C). Anal. Calcd. for C₃₃H₁₉N₂O₃: C, 80.47%; H, 4.09%; N, 5.69%; O, 9.75%. Found: C, 80.08%; H, 4.57%; N, 3.69%; O, 10.34%.

Example 3

Polymerization of AB₂ monomer in PPA/P₂O₅ at 160°C

15 Into a 100 mL resin flask equipped with a high torque mechanical stirrer, nitrogen inlet and outlet, a pressure regulator and an addition port, PPA (83% assay, 40g) was placed and stirred with dried nitrogen purging at 100°C for 10h. The monomer, 2,3-bis(4-phenyloxyphenyl)-6-quinoxaline-carboxylic acid (1.80g, 3.5 mmol) and P₂O₅ (10.0g) were added and the mixture heated to 130°C until it became a homogeneous 20 mixture. It took about 3h. The mixture was then heated at 160°C and it soon stuck to the stirring rod, rendering further efficient stirring/mixing impossible. It took around 3h. At the end of the reaction, water was added into the reaction vessel. The resulting lumps were broken up with the aid of a Waring blender, and the polymer product was

collected by suction filtration, washed with diluted ammonium hydroxide and then with a large amount of water, stirred in boiling water for 100h, and finally dried in the presence of phosphorous pentoxide under reduced pressure (1 mmHg) at 200°C for 48h. The yield was essentially quantitative (>99% yield). The polymer was insoluble in most
5 organic solvents and only formed gel in methanesulfonic acid (MSA). Anal. Calcd. for C₃₃H₁₉N₂O₃: C, 80.47%; H, 4.09%; N, 5.69%. Found: C, 78.34%; H, 4.34%; N, 5.27%.

Example 4

Polymerization in 1:10 w/w mixture of P₂O₅/Methanesulfonic acid (PPMA) at 110°C
Into a 100 mL 3-necked round bottom flask equipped with a mechanical stirrer,
10 nitrogen inlet and outlet, the monomer 2,3-bis(4-phenoxyphenyl)-6-quinoxaline-
carboxylic acid (1.0g, 1.96 mmol) and PPMA (10 mL) were added and heated to 110°C
for 8h. The color of mixture became dark purple. The mixture was poured into ice
water. The resulting purple precipitates were collected by suction filtration, washed with
diluted ammonium hydroxide and then with large amount of water, stirred in boiling
15 water for 48h, and finally dried in the presence of phosphorous pentoxide under
reduced pressure (1 mmHg) at 200°C for 48h. The yield was essentially quantitative
(>99% yield). Its intrinsic viscosity was 0.50 dL/g (MSA, 30 ± 0.1°C). Anal. Calcd. for
C₃₃H₁₉N₂O₃: C, 80.47%; H, 4.09%; N, 5.69%; O, 9.75%. Found: C, 78.12%; H, 4.33%;
N, 5.27%; O, 10.77%.

20 The polymerization results are summarized in Table 1.

Table 1. Polymerization Conditions and Yields of Polymer

Example	Media	P ₂ O ₅ (wt%) ^a	Temp. (°C)	Time (h)	[η] dL/g ^b	Yield (%)
1	PPA	0	130	48	0.07	~97
2	PPA	25	130	24	0.56	>99
3	PPA	25	160	3	Gel	>99
4	MSA	10	110	8	0.50	>99

(a) Relative to the amount of PPA used.

(b) Intrinsic viscosity measured in methanesulfonic acid (MSA) at 30±0.1°C.

Thermal Properties. The T_g's of the polymers were determined by DSC. The 5 thermograms were obtained on powder samples after they had been heated to 400°C and cooled to 20°C with heating and cooling rate of 10°C/min. The T_g value was taken as the mid-point of the maximum baseline shift from the second run. The polymer samples (Examples 1-4), which were prepared from different reaction conditions, displayed T_g's at 149°C, 113°C, not detectable, 91°C, in that order. Thermogravimetric 10 analysis of these polymers showed that they were heat-resistant; temperatures at which a 5% weight loss was observed were in the range of 505-525°C in air and 515-536°C in helium, respectively.

Example 5

Preparation of 3,5-dihydroxy-terminated polyetherketonequinoxaline

15 Into a 250 mL resin flask equipped with a high torque mechanical stirrer, nitrogen inlet and outlet, a pressure regulator and an addition port, PPA (83% assay, 80g), P₂O₅ (20.0g; 25 wt% relative to PPA used) were placed and heated to 130°C until it became homogeneous. After the reaction medium had been allowed to cool to 100°C, the monomer, 2,3-bis(4-phenoxyphenyl)-6-carboxyquinoxaline (4.0g, 3.92 mmol) and the 20 end-capper 3,5-dihydroxybenzoic acid (1.21g, 7.84 mmol) were added at the same time. The color of mixture became deep purple. The mixture was heated at 130°C for

2h, and at 160°C for 24h. When the polymerization was terminated, the mixture was moderately viscous. It was allowed to cool to 100°C (still fluid) to facilitate pouring into water. The resulting precipitates were collected by suction filtration, washed with diluted ammonium hydroxide and then with a large amounts of water, Soxhlet-extracted with
5 water for 50h and then with methanol for additional 50h, and finally dried in the presence of phosphorous pentoxide under reduced pressure (1 mmHg) at 100°C for 48h. The yield was essentially quantitative (>99% yield). The reduced viscosities at 0.5 g/dL were 0.09 dL/g (NMP, 30 ± 0.1°C) and 0.21 dL/g (MSA, 30 ± 0.1°C). Anal. Calcd. for C₄₀H₂₄N₂O₆: C, 76.43%; H, 3.85%. Found: C, 76.05%; H, 4.28%.

10

Example 6

Preparation of 3,5-diallyl-terminated polyetherketonequinoxaline

Into a 100 mL three-necked, round-bottomed flask equipped with a magnetic stirrer, nitrogen inlet, a condenser, and an addition port, hydroxyl-terminated hyperbranched polymer (example 5; 3.5 g, 5.57 mmol), potassium carbonate (4.0 g, 15 28.90 mmol), allyl bromide (3.37 g, 27.86 mmol), and dimethylacetamide (50 mL) were placed. The reaction mixture was then heated and maintained at 90-100°C for 16 h. During this time period, the orange solution became light yellow in color and homogeneous. After it had been allowed to cool down on its own, the mixture was filtered through a cake of Celite 545 to remove any insoluble salts. The filtrate was 20 poured into a beaker containing 5% hydrochloric acid (300 mL) and the mixture was warmed up to around 60-70°C for 2 h. The white powder was collected, air-dried, dissolved in dichloromethane, precipitated in methanol, collected, and dried under the reduced pressure in the presence of phosphorus pentoxide at 50°C for 48 h. The yield

was essentially quantitative. $[\eta] = 0.13 \text{ dL/g}$. Anal. Calcd. for $\text{C}_{46}\text{H}_{32}\text{N}_2\text{O}_6$: C, 77.95%; H, 4.55%; N, 3.95%; O, 13.54%. Found: C, 77.82%; H, 4.49%; N, 3.67%; O, 13.45%.

Having thus described exemplary embodiments of the present invention, it should be noted by those skilled in the art that the disclosures herein are exemplary only and that alternatives, adaptations and modifications may be made within the scope of the present invention.